INFORMATION DISCLOSURE STATEMENT	Atty. Docket No.: 232.00010120	Serial No.: 08/981,998
	Applicant(s): Pulst	
	Filing Date: May 11, 1998	Group: ~~1801~~ 1655

U.S. PATENT DOCUMENTS

Examiner Initial		Document Number	Date	Name	Class	SubClass	Filing Date If Appropriate
		5,552,282	09/03/96	Caskey et al.	435	6	
		5,650,270	07/22/97	Giese et al.	435	6	
		5,650,277	07/22/97	Navot et al.	435	6	
		5,741,645	04/21/98	Orr et al.	435	6	

FOREIGN PATENT DOCUMENTS

		Document Number	Date	Country	Class	SubClass	Translation Yes	Translation No
		WO 95/01437	01/12/95	PCT				
		WO 97/17445	05/15/97	PCT				X

OTHER DOCUMENTS (Including Authors, Title, Date, Pertinent Papers, etc.)

			Banfi, et al., "Identification and characterization of the gene causing type 1 spinocerebellar ataxia," Nature Genetics, 7, 513-519 (1994).
			Belal et al., "Clinical and genetic analysis of a Tunisian family with autosomal dominant cerebellar ataxia type 1 linked to the SCA2 locus," Neurology, 44, 1423-1426 (1994).
			Brook, "Retreat of the triplet repeat," Nat. Genet., 3, 279-281 (1993).
			Brunner et al., "Brief Report: Reverse Mutation In Myotonic Dystrophy," New Engl. J. Med., 328, 476-480 (1993).
			Filla et al., "Prevalence of hereditary ataxias and spastic paraplegias in Molise, a region of Italy," J. Neurol., 239, 351-353 (1992).
			Gispert et al., "Chromosomal assignment of the second locus for autosomal dominant cerebellar ataxia (SCA2) to chromosome 12q23-24.1," Nat. Genet., 4, 294-299 (1993).
			Imbert, "Cloning of the gene for spinocerebellar ataxia 2 reveals a locus with high sensitivity to expand CAG/glutamine repeats," Nature Genetics, 14, 285-291(1996).

EXAMINER Janine Enewold	Date Considered Jan 31, 2000

*Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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	Applicant(s): Pulst	
	Filing Date: May 11, 1998	Group: 1801

JE			Ioannou et al., "A new bacteriophage P1-derived vector for the propagation of large human DNA fragments," Nat. Genet., 6, 84-89 (1994).
JE			Kawaguchi et al., "CAG expansions in a novel gene for Machado-Joseph disease at chromosome 14q32.1," Nat. Genet., 8, 221-227 (1994).
JE			Koide et al., "Unstable expansion of CAG repeat in hereditary dentatorubral-pallidoluysian atrophy (DRPLA)," Nat. Genet., 6, 9-13 (1994).
JE			Kremer, et al., "Mapping of DNA Instability at the Fragile X to a Trinucleotide Repeat Sequence p(CCG)n," Science, 252, 1711-1714 (1991).
JE			Lopes-Cendes et al., "Confirmation of the SCA-2 Locus as an Alternative Locus for Dominantly Inherited Spinocerebellar Ataxias and Refinement of the Candidate Region," Am. J. Hum. Genet., 54, 774-781 (1994).
JE			MacDonald et al., "A Novel Gene Containing a Trinucleotide Repeat That Is Expanded and Unstable on Huntington's Disease Chromosomes," Cell, 72, 971-983 (1993).
JE			Mahadevan, et al., "Myotonic Dystrohpy Mutation: An Unstable CTG Repeat in the 3' Untranslated Region of the Gene," Science, 255, 1253-1255 (1992).
JE			Mandel, "Questions of expansion," Nat. Genet., 4, 8-9 (1993).
JE			Nagafuchi et al., "Dentatorubral and pallidoluysian atrophy expansion of an unstable CAG trinucleotide on chromosome 12p,"Nat. Genet., 6, 14-18 (1994).
JE			Orr et al., "Expansion of an unstable trinucleotide CAG repeat in spinocerebellar ataxia type 1," Nat. Genet., 4, 221-226 (1993).
JE			Polo et al., "Heredetiary Ataxias and Paraplegias in Cantabria, Spain," Brain, 114, 855-866 (1991).
JE			Pulst et al., "Anticipation in spinocerebellar ataxia type 2," Nat. Genet., 5, 8-10 (1993).
JE			Pulst et al., "Genetic and Physical Map of the Spinocerebellar Ataxia 2 (SCA2) Region on Human Chromosome 12," Neurology, 45, A422 (1995).
JE			Pulst et al., "Moderate expansion of a normally biallelic trinucleotide repeat in spinocerebellar ataxia type 2," Nature Genetics, 14, 269-276 (1996).
JE			Ranum et al., "Spinocerebellar ataxia type 5 in a family descended from the grandparents of President Lincoln maps to chromosome 11," Nat. Genet., 8, 280-284 (1994).

EXAMINER	Date Considered
Joanne Enewold	Jan 31, 2000

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	Applicant(s): Pulst	
	Filing Date: May 11, 1998	Group: 1801

JE				Rubensztein, et al., Phenotypic Characterization of Individuals with 30-40 CAG Repeats in the Huntington Disease (HD) Gene Reveals HD Cases with 36 Repeats and Apparently Normal Elderly Individuals with 36-39 Repeats," Am. J. Hum. Genet., 59, 16-22 (1996).
JE				Sanpei et al., "Identification of the spinocerebellar ataxia type 2 gene using a direct identification of repeat expansion and cloning technique, DIRECT," Nature Genetics, 14, 277-284 (1996).
JE				Stevanin et al., "Screening for proteins with polyglutamine expansions in autosomal dominant cerebellar ataxias," Human Molecular Genetics, 5, 1887-1892 (1996).
JE				Takiyama et al., "The gene for Machado-Joseph disease maps to human chromosome 14q," Nat. Genet., 4, 300-304 (1993).
JE				The WashU-Merck EST Project, "Soares parathyroid tumor NbHPA Homo sapiens cDNA clone," Accession No. W39162, May 15, 1996.
JE				Trottier et al., "Polyglutamine expansions as a pathological epitope in Huntington's disease and four dominant cerebellar ataxias," Letters to Nature, 378, 403-406 (1995).

EXAMINER	Date Considered
Janine Emerald	Jan 31, 2000

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